Response to Restriction Requirement

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Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the

application.

Claim 1 (Currently amended): A pharmaceutical composition comprising one compound,

which wherein the compound is a serotonin reuptake inhibitor, and another a second compound,

which wherein the second compound is a H₃ receptor antagonist, inverse agonist or partial

agonist having an affinity for the H₃ receptor below 0.5 μM.

Claim 2 (Currently amended): A pharmaceutical composition comprising a compound that is both

a H₃ receptor antagonist, inverse agonist or partial agonist having an affinity for the H₃ receptor

below $0.5 \mu M$, and a serotonin reuptake inhibitor.

Claim 3 (Currently amended): A method of augmenting and/or providing faster onset of

the therapeutic effect of a serotonin reuptake inhibitor, comprising administering to a patient in

need thereof a therapeutically effective amount of a H₃ receptor antagonist, inverse agonist or

partial agonist having an affinity for the H₃ receptor below 0.5 µM.

Claim 4 (Currently amended): A method of treating depression or an affective disorder,

comprising administering a therapeutically effective amount of a H₃ receptor antagonist, inverse

agonist or partial agonist having an affinity for the H₃ receptor below 0.5 µM to a patient being

treated with a serotonin reuptake inhibitor and in need thereof.

Claim 5 (Previously presented):

A method of treating depression or an

affective disorder, comprising administering to a patient in need thereof a therapeutically

effective amount of a pharmaceutical composition according to claim 1.

Claim 6 (Currently amended): The method of claim 5, wherein the compound serotonin

reuptake inhibitor is a selective serotonin reuptake inhibitor.

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Claim 7 (Currently amended): The method of claim 5, wherein the compound H₃ receptor antagonist, inverse agonist or partial agonist is selective for the H₃ receptor.

Claim 8 (Currently amended): The method of claim 5, wherein the <u>second</u> compound is an antagonist or an inverse agonist at the H₃ receptor.

Claim 9 (Currently amended): The method of claim $\underline{5}$ [[8]], wherein the <u>second</u> compound is a H_3 receptor antagonist.

Claim 10 (Currently amended): The method of claim [[6]] 5, wherein the serotonin reuptake inhibitor is selected from citalopram, escitalopram, fluoxetine, sertraline, paroxetine, fluoxamine, venlafaxine, dapoxetine, duloxetine, vilazodone, nefazodone, imipramine, femoxetine, or [[and]] clomipramine.

Claim 11 (Currently amended): The method of claim 5, wherein the H3 receptor ligand H3 receptor antagonist, inverse agonist or partial agonist is selected from Thioperamide, Ciproxifan, Iodophenpropit, GR 175737, Iodoproxyfan, Proxifan, Perceptin, JB 98064, VUF 9153, A 304121, ABT923, ABT 834, A 923, A 320436, A 331440, A 349413, A 349821, A 417022, A 423579, A 424835, A 431404, AQ 0145, FUB 181, FUB 360, FUB 407, FUB 637, FUB 836, GR 168320, GSK 189254A, GSK 207040A, GT 2016, GT 2104, GT 2209, GT 2212, GT 2227, GT 2232, GT 2390, GT 2349, GT 2355, GT 2394, Imoproxifan, Impentamine, JNJ 5207852, NNC 0038 0000 1049, NNC 0038 0000 1202, SCH 50971, SCH 79687, UCL 1199, UCL 1283, UCL 1390, UCL 1409, UCL 1860, UCL 1972, UCL 2065, UCL 2138, UCL 2173, UCL 2283, Verongamine, VUF 4163, VUF 5000, or [[and]] VUF 5182.

Claim 12 (Previously presented): The pharmaceutical composition according to claim 1, further comprising a pharmaceutically acceptable carrier or diluent.

Claim 13 (Currently amended): The pharmaceutical composition according to claim 1, wherein the serotonin reuptake inhibitor used is a selective serotonin reuptake inhibitor.

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Claim 14 (Previously presented): The pharmaceutical composition according to claim 1, wherein the H₃ antagonist, inverse agonist or partial agonist is selective for the H₃ receptor.

Claim 15 (Currently amended): The pharmaceutical composition according to claim 1, wherein the second compound H_3 ligand is a H_3 receptor antagonist.

Claim 16 (Currently amended): The pharmaceutical composition according to claim 1, wherein the serotonin uptake inhibitor is selected from citalopram, escitalopram, fluoxetine, sertraline, paroxetine, fluoxamine, venlafaxine, dapoxetine, duloxetine, vilazodone, nefazodone, imipramine, femoxetine, or [[and]] clomipramine.

Claim 17 (Currently amended): The pharmaceutical composition according to claim 1, wherein the H₃ ligand antagonist, inverse agonist or partial agonist is selected from from Thioperamide, Ciproxifan, Iodophenpropit, GR 175737, Iodoproxyfan, Proxifan, Perceptin, JB 98064, VUF 9153, A 304121, ABT923, ABT 834, A 923, A 320436, A 331440, A 349413, A 349821, A 417022, A 423579, A 424835, A 431404, AQ 0145, FUB 181, FUB 360, FUB 407, FUB 637, FUB 836, GR 168320, GSK 189254A, GSK 207040A, GT 2016, GT 2104, GT 2209, GT 2212, GT 2227, GT 2232, GT 2390, GT 2349, GT 2355, GT 2394, Imoproxifan, Impentamine, JNJ 5207852, NNC 0038 0000 1049, NNC 0038 0000 1202, SCH 50971, SCH 79687, UCL 1199, UCL 1283, UCL 1390, UCL 1409, UCL 1860, UCL 1972, UCL 2065, UCL 2138, UCL 2173, UCL 2283, Verongamine, VUF 4163, VUF 5000, or [[and]] VUF 5182.

Claim 18 (Currently amended): The method of claim 5, wherein the active ingredients serotonin uptake inhibitor and H₃ antagonist, inverse agonist or partial agonist are administered by simultaneous administration.

Claim 19 (Currently amended): The method of claim 5, wherein the active ingredients serotonin uptake inhibitor and H₃ antagonist, inverse agonist or partial agonist are administered in the same unit dosage form.

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Claim 20 (Currently amended): The method of claim 5, wherein the active ingredients serotonin uptake inhibitor and H₃ antagonist, inverse agonist or partial agonist are administered

by sequential administration.

Claim 21 (Currently amended): The method of claim 5, wherein the active ingredients

serotonin uptake inhibitor and H₃ antagonist, inverse agonist or partial agonist are administered

in discrete dosage forms.

Claim 22 (Currently amended): A method for identifying compounds useful for the

treatment of depression or an affective disorder, comprising, in any order:

(a) measuring the ability of test compounds to inhibit serotonin reuptake and selecting

the compounds that have an IC50 value below 50 nM;

(b) measuring the affinity of test selected compounds to the H₃ receptor and further

selecting the compounds that have an affinity for the H_3 receptor below 0.5 μ M,

and thereafter measuring the efficacy of the selected compounds at the H₃ receptor and

further selecting the compounds which are antagonists, inverse agonists or partial agonists at the

H₃ receptor.

Claim 23 (Previously presented):

The method according to claim 22 wherein the

compound has an affinity in step (b) of less than 50 nM.

Claim 24 (Previously presented):

The method according to claim 23, wherein the

compound has an affinity in step (b) of less than 10 nM.

Claim 25 (Currently amended):

A compound that inhibits serotonin reuptake and

has an IC₅₀ value below 50 nM; and has an affinity to the H₃ receptor below 0.5 μM.

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Claim 26 (Previously presented): A method of treating depression or an affective disorder, comprising administering to a patient in need thereof a therapeutically effective amount of a pharmaceutical composition according to claim 2.

Claim 27 (Currently amended): The method of claim 26, wherein the <u>compound</u> serotonin reuptake inhibitor is a selective serotonin reuptake inhibitor.

Claim 28 (Currently amended): The method of claim 26, wherein the <u>compound</u> H₃ receptor antagonist, inverse agonist or partial agonist is selective for the H₃ receptor.

Claim 29 (Currently amended): The method of claim 26, wherein the <u>compound H₃</u> receptor antagonist, inverse agonist or partial agonist is an antagonist or an inverse agonist at the H₃ receptor.

Claim 30 (Currently amended): The method of claim 29, wherein the <u>compound</u> H₃ receptor antagonist, inverse agonist or partial agonist is a H₃ receptor antagonist.

Claim 31 (Previously presented): The pharmaceutical composition according to claim 2, further comprising a pharmaceutically acceptable carrier or diluent.

Claim 32 (Currently amended): The pharmaceutical composition according to claim 2, wherein the <u>compound</u> H_3 -ligand is a H_3 receptor antagonist.

Claim 33 (Currently amended): The method of claim 4, wherein the affective disorder is selected from an anxiety disorder disorders, generalized anxiety disorder, panic anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit hyperactivity disorder, or drug abuse or any other disorder responsive to a serotonin reuptake inhibitor.

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Claim 34 (Currently amended): The method of claim 33, wherein the eating disorder is

selected from, bulimia, anorexia, or [[and]] obesity.

Claim 35 (Currently amended): The method of claim 5, wherein the affective disorder is

selected from an anxiety disorder disorders, generalized anxiety disorder, panic anxiety,

obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social

anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual

syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit

hyperactivity disorder, or drug abuse or any other disorder responsive to a serotonin reuptake

inhibitor.

Claim 36 (Currently amended): The method of claim 35, wherein the eating disorder is

selected from, bulimia, anorexia, or [[and]] obesity.

Claim 37 (Currently amended): The method of claim 22, wherein the affective disorder is

selected from an anxiety disorder disorders, generalized anxiety disorder, panic anxiety,

obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social

anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual

syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit

hyperactivity disorder, or drug abuse or any other disorder responsive to a serotonin reuptake

inhibitor.

Claim 38 (Currently amended): The method of claim 37, wherein the eating disorder is

selected from, bulimia, anorexia, or [[and]] obesity.

Claim 39 (Currently amended): The method of claim 26, wherein the affective

disorder is selected from an anxiety disorder disorders, generalized anxiety disorder, panic

anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder,

social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual

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syndrome, <u>a</u> cognitive <u>disorder</u> <u>disorders</u>, <u>an</u> impulse control <u>disorder</u> <u>disorders</u>, attention deficit hyperactivity disorder, <u>or</u> drug abuse <u>or any other disorder responsive to a serotonin reuptake inhibitor</u>.

Claim 40 (Currently amended): The method of claim 39, wherein the eating disorder is selected from, bulimia, anorexia, or [[and]] obesity.

Claim 41 (Previously presented): The compound of claim 25, wherein the compound has an affinity to the H_3 receptor of less than 50 nM.

Claim 42 (Previously presented): The compound of claim 41, wherein the compound has an affinity to the H₃ receptor of less than 10 nM.